



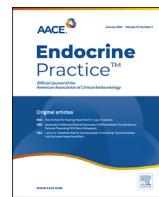
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## Review Article

## Vitamin D and Its Potential Benefit for the COVID-19 Pandemic

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## ABSTRACT

Vitamin D is known not only for its importance for bone health but also for its biologic activities on many other organ systems. This is due to the presence of the vitamin D receptor in various types of cells and tissues, including the skin, skeletal muscle, adipose tissue, endocrine pancreas, immune cells, and blood vessels. Experimental studies have shown that vitamin D exerts several actions that are thought to be protective against coronavirus disease (COVID-19) infectivity and severity. These include the immunomodulatory effects on the innate and adaptive immune systems, the regulatory effects on the renin-angiotensin-aldosterone-system in the kidneys and the lungs, and the protective effects against endothelial dysfunction and thrombosis. Prior to the COVID-19 pandemic, studies have shown that vitamin D supplementation is beneficial in protecting against risk of acquiring acute respiratory viral infection and may improve outcomes in sepsis and critically ill patients. There are a growing number of data connecting COVID-19 infectivity and severity with vitamin D status, suggesting a potential benefit of vitamin D supplementation for primary prevention or as an adjunctive treatment of COVID-19. Although the results from most ongoing randomized clinical trials aiming to prove the benefit of vitamin D supplementation for these purposes are still pending, there is no downside to increasing vitamin D intake and having sensible sunlight exposure to maintain serum 25-hydroxyvitamin D at a level of least 30 ng/mL (75 nmol/L) and preferably 40 to 60 ng/mL (100–150 nmol/L) to minimize the risk of COVID-19 infection and its severity.

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## Introduction

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is the new strain of coronavirus that causes coronavirus disease 2019 (COVID-19).<sup>1,2</sup> Due to the high infectivity and transmissibility of this novel virus, COVID-19 quickly became a global pandemic that has already affected at least 219 countries since its emergence from Wuhan, China in December 2019.<sup>2,3</sup> The most common clinical manifestations of COVID-19 include fever, fatigue, anorexia, myalgia, cough, sputum production, and dyspnea.<sup>4,5</sup> Although the

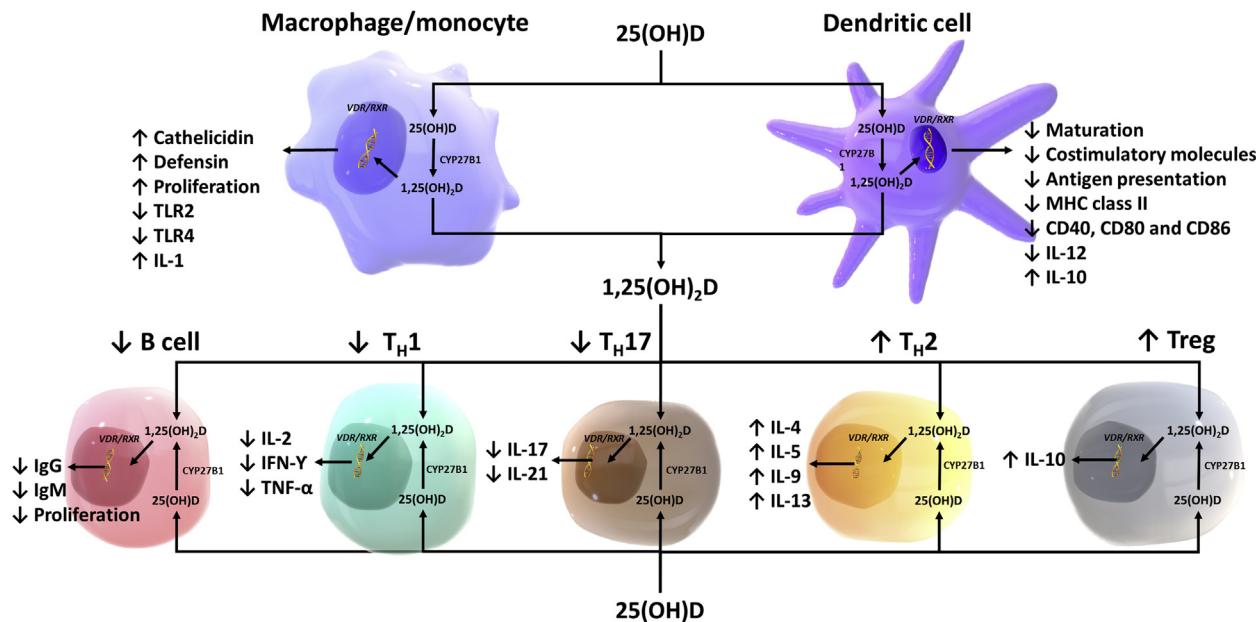
majority of patients with COVID-19 are either asymptomatic or develop only mild respiratory symptoms, a significant number of patients develop severe complications that result in morbidity and mortality, including acute respiratory distress syndrome (ARDS), arterial and venous thrombosis, multi-organ failure, and septic shock, among others.<sup>4,5</sup> Factors known to be associated with increased susceptibility to severe outcomes are advanced age, cancer, immunocompromised state, chronic kidney disease, chronic respiratory disease, cardio-metabolic disorders and smoking.<sup>6</sup> The elderly, African Americans, patients with obesity, and nursing home residents<sup>7,8</sup> have disproportionately higher rates of infection, morbidity, and mortality from COVID-19. These populations are also known as being at high risk for vitamin D deficiency.<sup>9–12</sup> Thus, vitamin D deficiency could potentially contribute to higher COVID-19 positivity, morbidity, and mortality rates appreciated in these populations.

Vitamin D is not only known for its importance for bone health but is also recognized for its potential protective effects against multiple chronic diseases as well as its immunomodulatory

**Abbreviations:** ACE2, angiotensin converting enzyme 2; ARDS, acute respiratory distress syndrome; COVID-19, coronavirus disease; IL, interleukin; IU, international units; OR, odds ratio; PBMC, peripheral blood mononuclear cell; RAAS, renin-angiotensin-aldosterone; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2; T<sub>H</sub>1, T helper 1; T<sub>H</sub>17, T helper 17; VDR, vitamin D receptor.

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**Fig. 1.** Schematic representation of paracrine and intracrine function of vitamin D and its metabolites and actions of 1,25-dihydroxyvitamin D on the innate and adaptive immune systems. 1,25(OH)<sub>2</sub>D = 1,25-dihydroxyvitamin D; 25(OH)D = 25-hydroxyvitamin D; IFN-γ = interferon-γ; IL = interleukin; MHC = membrane histocompatibility complex; T<sub>H</sub>1 = T helper 1; T<sub>H</sub>2 = T helper 2; T<sub>H</sub>17 = T helper 17; T<sub>reg</sub> = regulatory T cell; TLR2 = toll-like receptor 2; TLR4 = toll-like receptor 4; TNF-α = tumor necrosis factor-α. Reproduced with permission from Holick, 2020.

activities.<sup>10,11,13</sup> With the global prevalence of vitamin D deficiency (defined by serum 25-hydroxyvitamin D [25(OH)D] level of <20 ng/mL) and insufficiency (defined by serum 25(OH)D level of 20–<30 ng/mL) of 40% to 100%,<sup>14–17</sup> correcting vitamin D deficiency would be a cost-effective intervention to alleviate the burden of this pandemic at a populational level. The aim of this review is to discuss potential biological mechanisms by which vitamin D could be protective against COVID-19 and to summarize evidence from observational studies and clinical trials that have demonstrated the direct and indirect links between vitamin D and COVID-19.

## Sources, Synthesis, and Metabolism of Vitamin D

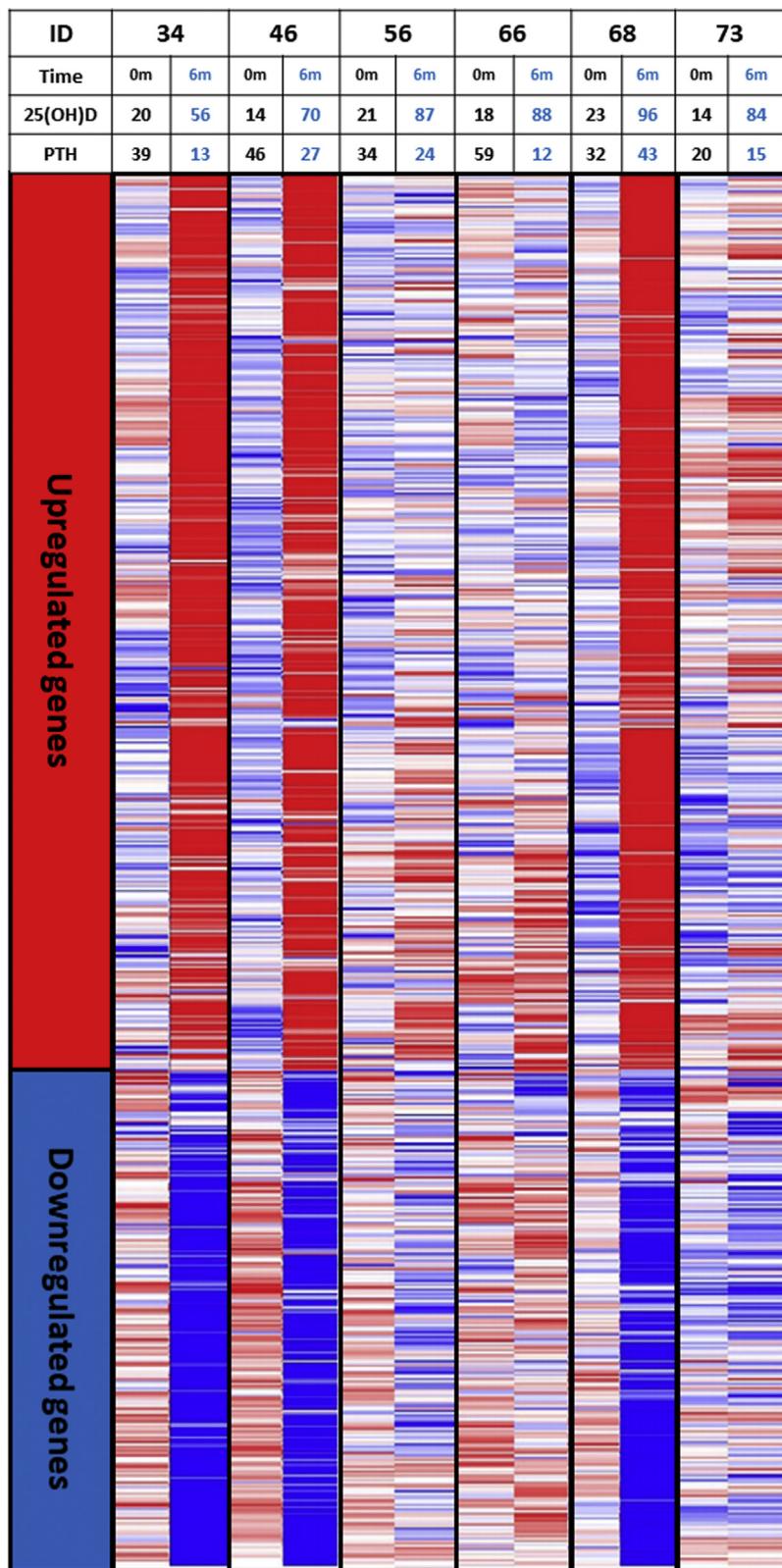
Vitamin D is responsible for regulating calcium and phosphate metabolism and maintaining a healthy mineralized skeleton. It is also known for its biologic activities on various types of tissues including the immune system.<sup>10,11,13,18–20</sup> There are 2 forms of vitamin D: vitamin D<sub>2</sub> (ergocalciferol) and vitamin D<sub>3</sub> (cholecalciferol). Vitamin D<sub>2</sub>, synthesized from ergosterol, is found in sun dried and ultraviolet irradiated mushrooms and yeast, while vitamin D<sub>3</sub> is synthesized from endogenous 7-dehydrocholesterol in the skin and can be found naturally in oily fish and cod liver oil, as well as in meat in the form of 25(OH)D<sub>3</sub>.<sup>10,11,21–23</sup> Once entering the circulation, vitamin D (vitamin D<sub>2</sub> and D<sub>3</sub>) is converted by several vitamin D-25-hydroxylases (ie, CYP2R1, CYP27A1, CYP2C11, CYP2J3, CYP3A4) in the liver into 25(OH)D, the major circulating metabolite of vitamin D. 25(OH)D is then metabolized by the 25-hydroxyvitamin D-1 $\alpha$ -hydroxylase (CYP27B1) to the biologically active form, 1,25-dihydroxyvitamin D [1,25(OH)<sub>2</sub>D].<sup>24</sup> The kidneys are the main site of conversion of 25(OH)D into the circulating bioavailable 1,25(OH)<sub>2</sub>D, which is responsible for regulating intestinal calcium absorption and bone calcium mobilization.<sup>10,11</sup> Furthermore, CYP27B1 is expressed in several other tissues, including parathyroid glands, breast, colon, keratinocytes, microglia, and immune cells, where 1,25(OH)<sub>2</sub>D is produced and exerts its autocrine, paracrine, and intracrine functions by binding

with the intracellular vitamin D receptor (VDR), which subsequently leads to up- or down-regulation of a multitude of genes.<sup>10,11</sup>

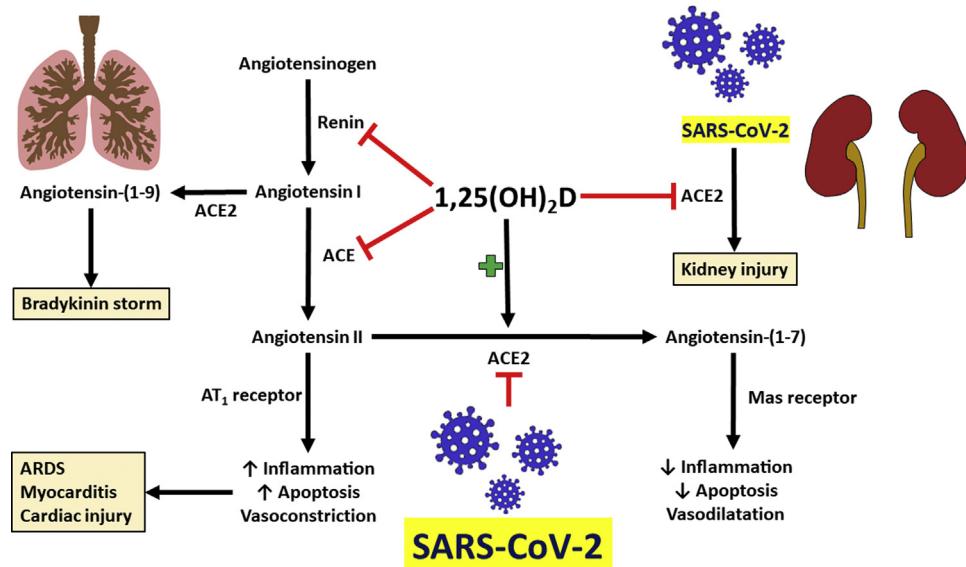
## Vitamin D and Immune Function

Due to the presence of the VDR in most tissues, including the skin, skeletal muscle, adipose tissue, endocrine pancreas, immune cells, and blood vessels, vitamin D has been shown to have a multitude of nonskeletal biological activities. In particular, vitamin D is considered an immunomodulatory agent that regulates both innate and adaptive immune systems (Fig. 1).<sup>10,11,13,18–20</sup> Activated macrophages express CYP27B1 that converts 25(OH)D into 1,25(OH)<sub>2</sub>D. 1,25(OH)<sub>2</sub>D, in turn, induces the macrophage production of the endogenous antimicrobial peptides, cathelicidins, and defensins.<sup>18,19,25</sup> Furthermore, 1,25(OH)<sub>2</sub>D has been shown to alter the activity of different types of T helper cells by promoting a shift from T helper 1 (T<sub>H</sub>1) and T helper 17 (T<sub>H</sub>17) to T helper 2 immune profile and facilitating differentiation of regulatory T cells.<sup>26–29</sup> In addition, both cytotoxic T lymphocytes and B cells, when activated, upregulate their VDR, suggesting a coordinated regulation of the VDR signaling pathway and response to stimuli of these components of the adaptive immune system.<sup>30–32</sup>

The effect of vitamin D supplementation on immune function has been well-demonstrated in a recent study that evaluated broad gene expression in peripheral blood mononuclear cells (PBMCs) after orally supplementing various doses of vitamin D.<sup>33–35</sup> Thirty healthy adults with vitamin D insufficiency (25(OH)D 20–<30 ng/mL or 50–<75 nmol/L) or deficiency (25(OH)D <20 ng/mL or <50 nmol/L) were randomized to receive 600, 4000, or 103000 international units (IU) per day of vitamin D<sub>3</sub> for 6 months and were found to have dose-dependent alteration in broad gene expression with 162, 320, and 1289 genes up- or down-regulated in their PBMCs, respectively.<sup>33</sup> Equally interesting if not more is that some individuals might respond to vitamin D more or less than others, as high interindividual difference in responsiveness to vitamin D supplementation has been observed (Fig. 2). In the same clinical trial, those who



**Fig. 2.** Heatmaps of vitamin D responsive genes whose expression response variation in 6 vitamin D-deficient subjects taking 10 000 international units per day of vitamin D3 for 6 months showed that 3 subjects had a robust response in gene expression compared to the other 3 subjects, who had minimum-to-modest responses even though these subjects raised their blood levels of 25(OH)D in the same range of ~60 to 90 ng/mL. 0m = 0 month; 6m = 6 months; 25(OH)D = 25-hydroxyvitamin D; PTH = parathyroid hormone. Reproduced with permission from Holick, 2019.



**Fig. 3.** Schematic representation of the effects of 1,25(OH)<sub>2</sub>D on the renin-angiotensin-aldosterone system. SARS-CoV-2 uses the ACE2 as the main receptor entry site and downregulates ACE2 in the lungs. This causes the accumulation of angiotensin II, which causes inflammation and apoptosis in the lungs and systemic vasoconstriction by interacting with the AT<sub>1</sub> receptor, leading to COVID-related complications including ARDS, myocarditis, and cardiac injury. 1,25(OH)<sub>2</sub>D inhibits renin and ACE and induces the expression of ACE2 in the lungs, thereby reducing the accumulation of angiotensin II. Inhibition of renin expression may also result in decreased flux of angiotensin I to angiotensin-(1-9), thereby mitigating bradykinin storm. Additionally, 1,25(OH)<sub>2</sub>D may inhibit ACE2 expression in the renal tubular cells, which is thought to be protective against COVID-associated kidney injury by reducing the viral direct cytopathic effects on the cell. 1,25(OH)<sub>2</sub>D = 1,25-dihydroxyvitamin D; ACE = angiotensin converting enzyme; ACE2 = angiotensin converting enzyme 2; ARDS = acute respiratory distress syndrome; AT<sub>1</sub> receptor = angiotensin II type 1 receptor; COVID-19 = coronavirus disease 2019; SARS-CoV-2 = severe acute respiratory distress syndrome coronavirus 2 (Copyright Holick, 2021).

received this same dose of vitamin D and raised their serum concentrations of 25(OH)D to the same degree showed marked differences in the level of expression of the same genes.<sup>33</sup> In addition, different patterns of serum metabolomic profile were also observed between the subjects with robust and minimum-to-modest genomic responses.<sup>33,34</sup> These observations support of the findings from a previous clinical trial that gave 3200 IU of vitamin D<sub>3</sub> per day to 71 prediabetic patients for 5 months and revealed robust changes in broad gene expression in PBMCs only in about half of the subjects despite comparable serum concentrations of 25(OH)D.<sup>35</sup>

#### Potential Protective Effects of Vitamin D Against COVID-19

There are multiple biological explanations by which vitamin D could potentially be protective against infectivity and severity from COVID-19. These include vitamin D's immune- and nonimmune-mediated actions on several tissues via both genomic and nongenomic pathways. First, 1,25(OH)<sub>2</sub>D enhances the innate immune system by inducing not only the macrophages but also the respiratory epithelial cells to produce the antimicrobial peptide, cathelicidin LL-37.<sup>36</sup> This antimicrobial peptide not only acts against invading bacteria and fungi by destabilizing their cell membranes, but also exhibits direct antiviral activities against respiratory viruses by altering viability of host target cells and disrupting their envelopes.<sup>37–39</sup> This mechanism is supported by the result of a pilot clinical trial that gave a single enteral dose of 400 000 IU of vitamin D<sub>3</sub> or placebo to patients with sepsis and demonstrated an increase in serum cathelicidin in the treatment group compared with the placebo group.<sup>40</sup> More interestingly, it has been recently demonstrated in an experimental study using surface plasmon resonance analysis that LL-37 competitively binds to SARS-CoV-2 S protein, which, in turn, inhibits viral binding to the receptor ACE2 and most likely prevents viral entry into the cell.<sup>41</sup> In addition, cathelicidins were shown to prevent lung damage associated with oxygen toxicity.<sup>42</sup>

The second mechanism is related to the immunomodulatory effects of vitamin D on the adaptive immune system. As discussed in the previous section, 1,25(OH)<sub>2</sub>D has been shown to downregulate the activities of T<sub>H</sub>1 and T<sub>H</sub>17 and promote differentiation of regulatory T cells.<sup>26–29</sup> This leads to a decrease in the production of proinflammatory cytokines, including interleukin (IL)-6, IL-8, IL-12, tumor necrosis factor- $\alpha$ , and IL-17,<sup>26–29</sup> thereby alleviating the cytokine storm syndrome in patients with COVID-19 with high inflammatory burden and therefore preventing multi-organ dysfunction. Interestingly, vitamin D has also been shown to upregulate the expression of IL-10, which is thought to be a potential treatment target for COVID-19.<sup>43–46</sup> These potential immunologic effects of vitamin D are supported by multiple studies that reported the impact of vitamin D supplementation on reduction of inflammatory burden in T<sub>H</sub>1- and/or T<sub>H</sub>17-mediated autoinflammatory diseases such as rheumatoid arthritis,<sup>47</sup> psoriasis,<sup>48,49</sup> multiple sclerosis,<sup>50</sup> and inflammatory bowel disease.<sup>51</sup> In addition, it has been suggested that activation of the VDR in the pulmonary stellate cells might play a role in suppressing inflammation and fibrotic changes in the lungs of patients with COVID-19.<sup>52</sup>

Third, 1,25(OH)<sub>2</sub>D has been shown to regulate the renin-angiotensin-aldosterone (RAAS) system (Fig. 3),<sup>53,54</sup> and the effects are thought to be different among tissues. In an animal model, oral administration of alfalcacidiol (1 $\alpha$ -hydroxyvitamin D) was shown to inhibit ACE2 expression, which is the main receptor entry of SARS-CoV-2, in the renal tubular cells.<sup>54,55</sup> Therefore, 1,25(OH)<sub>2</sub>D likely exerts the same biologic on the kidney and therefore may be protective against COVID-associated kidney injury by reducing viral entry into the cell. It has been shown that SARS-CoV-2 infection downregulates ACE2 in the lungs.<sup>56</sup> This causes accumulation of angiotensin II, which is believed to play a role in the development of ARDS, myocarditis, and cardiac injury, the major severe complications of COVID-19.<sup>56</sup> In the lipopolysaccharide-induced acute lung injury animal model, 1,25(OH)<sub>2</sub>D was shown to suppress renin,

angiotensin converting enzyme, and angiotensin II expression and increase ACE2 expression.<sup>57,58</sup> These effects could potentially reduce the accumulation of angiotensin II and therefore reduce the risk of ARDS and cardiac injury, especially in patients with COVID-19 who have pre-existing dysregulation of the RAAS system, such as those with underlying hypertension, heart failure, and renal insufficiency.<sup>59</sup> Additionally, a mechanistic model generated from gene expression data of cells in bronchoalveolar lavage fluid from patients with COVID-19 and controls suggested that the inhibitory effect of 1,25(OH)<sub>2</sub>D on renin expression may result in decreased flux of angiotensin I to angiotensin-(1-9).<sup>60</sup> This mechanism is thought to help mitigate bradykinin storm, which has been shown to underlie the multiple organ dysfunction in COVID-19.<sup>60</sup>

Another action of vitamin D is its pleiotropic effects against endothelial cell dysfunction and vascular thrombosis, which may mitigate vascular leakage secondary to systemic inflammatory response and prevent COVID-associated arterial and venous thrombosis.<sup>61–63</sup> It has been shown in the primary dermal human microvascular endothelial cell model that vitamin D<sub>3</sub>, 25(OH)D<sub>3</sub>, and 1,25(OH)<sub>2</sub>D<sub>3</sub> stabilized vascular endothelial membranes via a nongenomic pathway.<sup>61</sup> Additionally, vitamin D<sub>3</sub>, which normally circulates at about 100 times higher concentration than 1,25(OH)<sub>2</sub>D<sub>3</sub>, was at least 10 times more potent than 1,25(OH)<sub>2</sub>D<sub>3</sub> and more than 1000 times more potent than 25(OH)D<sub>3</sub> in stabilizing the endothelium.<sup>61</sup> Furthermore, it has been shown in a uremic rat model that paricalcitol (19-nor-1,25[OH]<sub>2</sub>D<sub>2</sub>) could prevent the development of endothelial intracellular gaps and reduce endothelial damage.<sup>62</sup> Finally, vitamin D is known to exert direct and indirect antithrombotic activities by controlling the expression of multiple genes involved in the coagulation pathway.<sup>63</sup>

Despite multiple mechanisms suggesting potential benefits of vitamin D for COVID-19, 1,25(OH)<sub>2</sub>D is known to inhibit plasma cell differentiation and reduce immunoglobulin production by B cells in the settings of autoimmune disorders.<sup>30,64,65</sup> It is still unclear whether this biologic action could dampen the production of neutralizing antibodies and be detrimental in the setting of response to COVID-19 infection or COVID-19 vaccine. Further studies are required to investigate this aspect of vitamin D actions.

## Pre-COVID Evidence From Clinical Studies

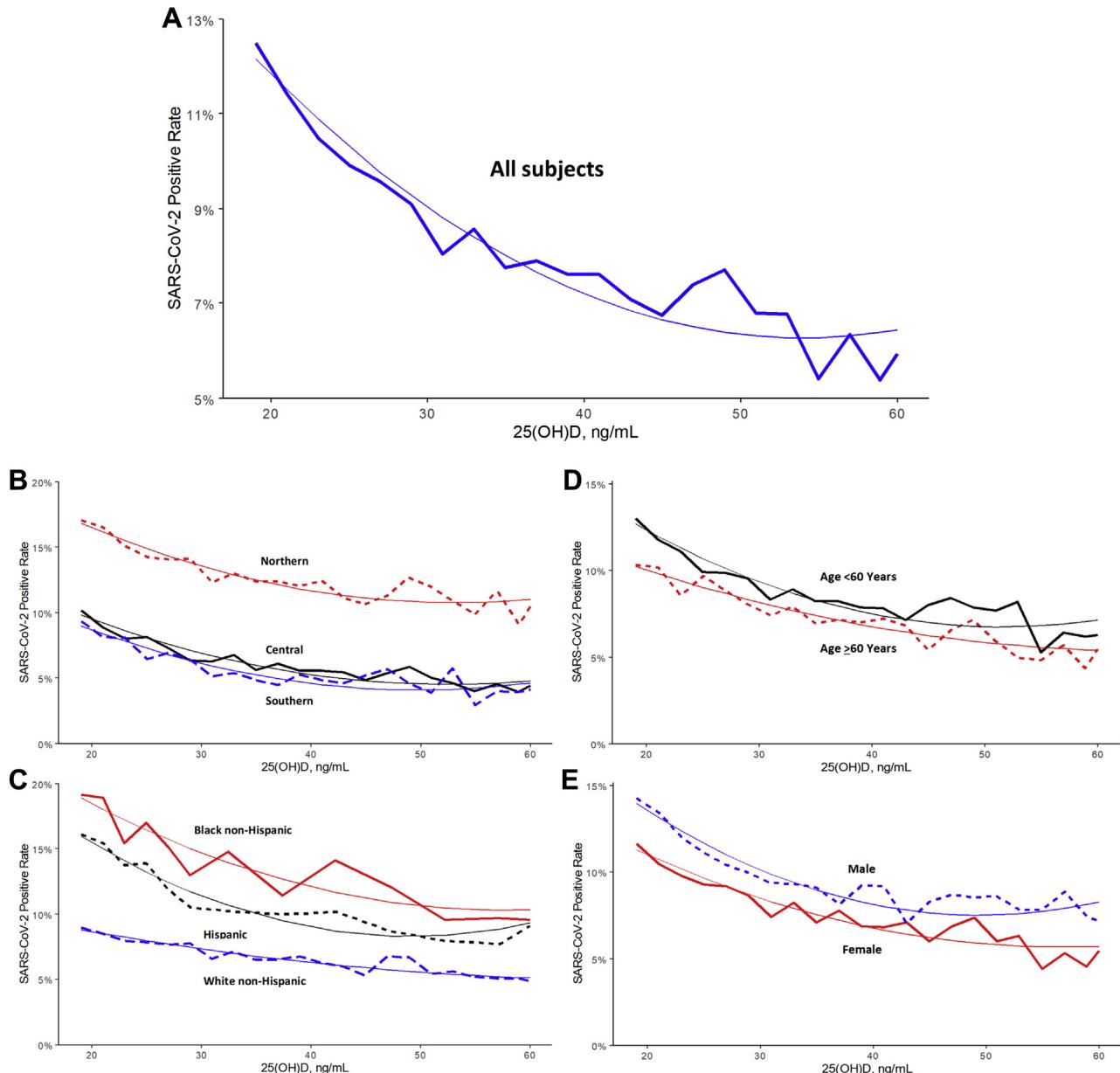
The outbreak of influenza infection is seasonal and usually occurs in the winter in high-latitude areas but is sporadic throughout the year in tropical areas.<sup>66,67</sup> The most likely explanation of this phenomenon is the seasonal variation of temperature, humidity, and intensity of ultraviolet radiation.<sup>68–70</sup> Another possible explanation for this outbreak pattern is the seasonal variation in serum concentrations of 25(OH)D, which reach the lowest levels in the winter.<sup>71</sup> This notion is supported by several studies that have shown the independent association between low concentration of serum 25(OH)D and incidence and severity of acute respiratory viral infection. For example, a cohort study in healthy adults demonstrated approximately 50% reduction in the risk of incident acute respiratory tract infection in those with serum 25(OH)D concentrations of  $\geq 38$  ng/mL (95 nmol/L).<sup>72</sup> A case-control study in 469 New Zealand children aged  $<2$  years demonstrated that those requiring hospitalization for acute respiratory infection had, significantly, 1.7-times higher odds of vitamin D deficiency than those with mild illnesses.<sup>73</sup> To illustrate the causal association, a randomized controlled trial gave 1200 IU of vitamin D<sub>3</sub> per day or placebo to 167 Japanese school children for 4 months and revealed that those who received vitamin D<sub>3</sub> supplementation had a significantly lower risk of influenza A infection compared with the placebo group (relative risk, 0.58; 95% CI, 0.34–0.99).<sup>74</sup> A more

recent meta-analysis of 25 randomized controlled trials showed that supplementation of vitamin D<sub>2</sub> or D<sub>3</sub> can protect against the development of acute respiratory tract infection compared with placebo (adjusted odds ratio [OR], 0.88; 95% CI, 0.81–0.96).<sup>75</sup> The protective effects were more pronounced in those with baseline 25(OH)D concentrations of less than 10 ng/mL or 25 nmol/L (adjusted OR, 0.30; 95% CI, 0.17–0.53).<sup>75</sup> It should, however, be noted that there was moderate statistical heterogeneity in this main meta-analysis, with the  $I^2$  value of 53.3%, and that most of the individual clinical trials included in the meta-analysis failed to demonstrate statistical significance of the impact of vitamin D supplementation.<sup>75</sup>

Prior to the COVID-era, sepsis was one of the major causes of morbidity and mortality among hospitalized patients in the intensive care unit.<sup>76</sup> A number of studies have shown the association between low concentrations of serum 25(OH)D and increased unfavorable outcomes in sepsis and critically ill patients.<sup>77,78</sup> However, the association between vitamin D status and sepsis outcomes might be bidirectional, as it is also probable that low serum 25(OH)D concentrations in patients with severe sepsis could be secondary to systemic inflammation that increases the activity of the 25(OH)D-24-hydroxylase that catabolizes 25(OH)D as well as causing extravascular leakage of the vitamin D-binding protein.<sup>79,80</sup> It should be noted that randomized clinical trials that investigated the impact of vitamin D supplementation on clinical outcomes of sepsis and critical illness have yielded mixed results. In a pilot study in 31 vitamin D-deficient patients who were on mechanical ventilation, administration of a single dose of enteral 500 000 or 250 000 IU of vitamin D<sub>3</sub> was found to decrease hospital length of stay compared with placebo.<sup>81</sup> In another randomized controlled trial that gave enteral 540 000 IU of vitamin D<sub>3</sub> followed by monthly maintenance doses of 90 000 IU for 5 months or placebo to 475 vitamin D-deficient critically ill patients, a significant decrease in-hospital mortality was observed in the subgroup of 200 patients with serum 25(OH)D  $< 12$  ng/mL or 30 nmol/L (hazard ratio, 0.56; 95% CI, 0.35–0.90).<sup>82</sup> On the other hand, in a larger clinical trial in 1360 patients with critical illness, administration of a single dose of enteral 540 000 IU of vitamin D<sub>3</sub> was not superior to placebo in reducing the risk of mortality and other clinical outcomes.<sup>83</sup> This negative result may suggest that it is too late for the critically ill patients to benefit from vitamin D supplementation and that vitamin D has to be given at the earlier stages of disease to demonstrate its survival benefit.<sup>84,85</sup>

## Current Evidence on Vitamin D and COVID-19

Multiple observational studies have reported the link between vitamin D status or serum 25(OH)D concentrations and risk of acquiring COVID-19 in many countries worldwide. For example, in a study using a national clinical laboratory database of the United States of 191 779 patients, SARS-CoV-2 positivity is strongly and inversely associated with circulating 25(OH)D concentrations, although the analysis was limited to 1 SARS-CoV-2 result per patient. The observed relationship was found to persist across latitudes, races, ethnicities, both sexes, and age ranges (Fig. 4).<sup>86</sup> This result is in line with that of a retrospective cohort study showing that deficient vitamin D status was associated with an increased risk of positive test for COVID-19 (relative risk, 1.77; 95% CI, 1.12–2.81) with likely sufficient vitamin D status after adjusting for potential confounders.<sup>87</sup> Another study in 50 hospitalized Korean patients with COVID-19 and 150 age- and sex-matched controls showed that the patients with COVID-19 were about 3 times more likely to be severely vitamin D-deficient (25[OH]D  $< 10$  ng/mL or 25 nmol/L) than the control group.<sup>88</sup> Another population-based study in 782 Israeli patients with COVID-19 and 7025 controls



**Fig. 4.** SARS-CoV-2 nucleic acid amplification test positivity rates and circulating 25(OH)D levels in all subjects (A) and stratified by latitude region (B), predominately Black non-Hispanic, Hispanic and White non-Hispanic zip codes (C), age group (D), and sex (E). Smooth lines represent the weighted second order polynomial regression fit to the data associating circulating 25(OH)D levels (x-axis) and SARS-CoV-2 positivity rates (y-axis). 25(OH)D = 25-hydroxyvitamin D; SARS-CoV-2 = severe acute respiratory distress syndrome coronavirus 2. (Copyright Kaufman, 2020<sup>86</sup> with permission.)

showed that vitamin D deficiency was independently associated with approximately 1.5-times higher odds of COVID-19 test positivity (adjusted OR, 1.50; 95% CI, 1.13–1.98).<sup>89</sup> In a study of 216 Spanish patients with COVID-19 and 197 population-based controls, vitamin D deficiency ( $25[\text{OH}]D < 20 \text{ ng/mL}$  or  $50 \text{ nmol/L}$ ) was found to be about 1.7-times more prevalent in COVID-19 cases than in the control group. Moreover, serum 25(OH)D concentrations were significantly lower in patients with COVID-19 after adjusting for potential confounders.<sup>90</sup> Nonetheless, a cohort study in 347 Italian hospitalized patients with positive and negative COVID-19 tests showed no association between vitamin D status and COVID-19 test positivity.<sup>90</sup> This negative finding is likely due to the fact that, unlike those of the other studies, hospitalized patients were recruited to be the control group.<sup>91</sup> A study using data from the United Kingdom biobank consisting of 348 598 participants

including 449 confirmed patients with COVID-19 reported that vitamin D was associated with COVID-19 infection univariately but not after adjustment for confounders. However, this study utilized serum concentrations of 25(OH)D measured during 2006 to 2000, which may not accurately reflect current vitamin D status.<sup>92</sup>

In addition to the promising data on the relationship between vitamin D status and risk of acquiring COVID-19, a growing amount of evidence from multiple observational studies has reported the connection between vitamin D status and risk of severity in patients with COVID-19. A meta-analysis of 27 studies reported that vitamin D deficiency in patients with COVID-19 was significantly associated with higher risks of severe infection (OR, 1.64; 95% CI, 1.30–2.09), hospitalization (OR, 1.81; 95% CI, 1.42–2.21), and mortality (OR, 1.92; 95% CI, 1.06–2.58).<sup>93</sup> Several more recent studies in many different regions worldwide have addressed the same

question with relatively inconsistent results.<sup>93–100</sup> This could be due to different patient characteristics and study design across the studies.

There are some issues that are worth noting while processing the evidence. First, vitamin D deficiency is associated with presence and disease burden of several comorbidities such as cardiovascular disorders, chronic kidney disease, and obesity.<sup>101–103</sup> Therefore, the observed association might be in part confounded by these factors, although most studies have already attempted to address this with multivariate analysis.<sup>98–100,104</sup> Second, the association between vitamin D status at the time of hospitalization and outcomes of acute inflammatory illness is likely due in part to reverse causation. A low level of serum 25(OH)D could also be secondary to systemic inflammatory response, which results in vascular leakage of vitamin D-binding protein and albumin as well as increased catabolism of 25(OH)D.<sup>105,106</sup> Third, vitamin D might benefit each individual differently as significant inter-individual difference in responsiveness to vitamin D supplement has been reported.<sup>33–35</sup> Additionally, aged individuals may benefit from vitamin D more than young adults as they tend to have higher inflammatory burden of COVID-19. This notion is supported by the observation in some studies that showed a stronger association between vitamin D status and COVID-19 severity in elderly patients.<sup>93,107</sup> Finally, some studies that reported positive association utilized previous laboratory data<sup>86,89,92</sup> and used the diagnostic code for vitamin D deficiency from the medical record database to define vitamin D status.<sup>98</sup> It is likely that an individual who was found to have vitamin D deficiency prior to the infection would have been treated for vitamin D deficiency and then became vitamin D repleted by the time they were infected. This indicates that there might be the legacy effect of being vitamin D-sufficient and that raising serum 25(OH)D concentrations over a short period of time might not be as effective as maintaining serum 25(OH)D concentrations in a preferred range of 40 to 60 ng/mL (100–150 nmol/L) over the long term.<sup>12</sup>

Given the promising evidence on the potential benefit of vitamin D against COVID-19, a number of ongoing randomized controlled trials have been conducted with the aim to investigate the impact of vitamin D supplementation of different forms and dosing regimens. A pilot randomized clinical trial gave oral 25(OH)D<sub>3</sub> (calcifediol) or placebo to 76 patients with COVID-19 and showed that the treatment group had a markedly reduced rate of intensive care unit admission (2% vs 50%,  $P < .001$ ).<sup>108</sup> However, in a larger randomized controlled trial giving 240 hospitalized patients with moderate-to-severe COVID-19 200 000 IU of vitamin D<sub>3</sub> or placebo, there were no differences in length of hospital stay, in-hospital mortality, admission to intensive care unit, or mechanical ventilation requirement.<sup>109</sup> This emphasizes that the immunomodulatory effects of vitamin D are likely to be the results of its long term rather than short-term actions.

### **Recommended Serum 25-Hydroxyvitamin D Concentrations to Help Fight the COVID 19 Pandemic**

It is largely controversial as to what concentration of serum 25(OH)D would provide optimal benefit for bone health, overall health benefits, and prevention against COVID-19. Serum 25(OH)D concentration of higher than 15 to 20 ng/mL (37.5–50 nmol/L) would be sufficient for prevention of rickets, osteomalacia, and symptomatic hypocalcemia.<sup>110</sup> Notably, hypocalcemia is shown to be highly prevalent and associated with hospitalization in patients with COVID-19. Whether and how much serum 25(OH)D would be protective against hypocalcemia in patients with COVID-19 requires further investigation.<sup>111</sup> However, it is recommended that serum

25(OH)D concentration should be above 30 ng/mL (75 nmol/L) to maximize the calcemic effects of vitamin D and minimize the risk of secondary hyperparathyroidism that predisposes to osteoporosis.<sup>12</sup> It is worth considering the historical evidence to postulate vitamin D status in our hunter-gatherer forefathers. Hadza tribesmen and Maasai herders were reported to have serum concentrations of 25(OH)D in the range of 40 to 60 ng/mL (100–150 nmol/L).<sup>9,112,113</sup> This range is in line with that reported not only in population-based studies to be associated with the lowest risk of chronic diseases and all-cause mortality<sup>9,11,113–116</sup> but also in recent studies to be associated with decreased risks of COVID-19 infection and its severity.<sup>86–90,93</sup> With minimal sunlight exposure, an adult would require ingestion of 4000 to 6000 IU of vitamin D<sub>3</sub> or vitamin D<sub>2</sub> daily to maintain serum 25(OH)D in the preferred range of 40 to 60 ng/mL (100–150 nmol/L).<sup>12</sup> Obese adults require 2 to 3 times more vitamin D to maintain the same serum concentrations of 25(OH)D.<sup>12,117</sup>

On average, approximately 40% and 60% of children and adults have circulating concentrations of 25(OH)D <20 ng/mL (50 nmol/L) and <30 ng/mL (75 nmol/L), respectively.<sup>116</sup> This already high prevalence of vitamin D deficiency/insufficiency tends to be further aggravated by the lack of sunlight exposure and outdoor activity as a result of the pandemic lockdown. Thus, patients hospitalized with COVID-19 are likely to be vitamin D-deficient or insufficient, and, therefore, it is reasonable to institute as standard of care to measure serum 25(OH)D level and to give at least 1 single dose of 80 000 to 100 000 IU of vitamin D to all vitamin D-deficient (25[OH]D <20 ng/mL or 50 nmol/L) or insufficient (25[OH]D 20–<30 ng/mL or 50–<75 nmol/L) patients with COVID-19 with a normal body mass index and at least 200 000 IU for those with obesity (body mass index >30 kg/m<sup>2</sup>) after being hospitalized.<sup>12,85,108</sup> It is noteworthy that optimal magnesium status may be important for optimizing vitamin D status.<sup>118,119</sup> Therefore, maintaining magnesium status by giving magnesium supplementation with high-dose vitamin D may benefit in this situation. Additionally, corticosteroids have become a mainstay treatment for COVID-19 in patients with high inflammatory burden. It should be noted that corticosteroids and some other medications (eg, antiepileptics and antiretrovirals) affect the steroid and xenobiotic receptor or the pregnane X receptor, causing increased catabolism of 25(OH)D and 1,25(OH)<sub>2</sub>D into inactive water-soluble carboxylic acid derivatives.<sup>12</sup> Thus, patients who receive any of these medications should also be given an increased dose of vitamin D of 200 000 IU.<sup>12</sup> Finally, if hospitalized more than 1 week, with minimal sunlight exposure and dietary intake of vitamin D, they should continue to receive the daily or the equivalent weekly dose of about 2000 to 5000 IU per day and 6000 to 10 000 IU per day for those with obesity or receiving corticosteroids.<sup>12</sup> This strategy is proposed to ensure serum 25(OH)D level of at least 30 ng/mL (75 nmol/L) throughout hospitalization. Further clinical trials are required to examine the clinical benefits or risks of this strategy specifically on COVID-19-related outcomes.

### **Conclusion**

Vitamin D is known not only for its importance for calcium and phosphate metabolism but also for its biologic actions on immune modulation. This is because of the presence of the VDR in most types of cells, especially the immune cells, including activated T and B lymphocytes and macrophages. Experimental studies have shown that vitamin D exerts several biological activities that are thought to be protective against COVID-19. These include the immunomodulatory effects on the innate and adaptive immune systems, the regulatory effects on the RAAS in the kidneys and the lungs, and the protective effects against endothelial dysfunction and thrombosis. Prior to the COVID-era, it was reported that

vitamin D supplementation is beneficial in protecting against risk of respiratory viral infection and may improve outcomes in sepsis and critically ill patients. There are a growing number of data suggesting the link between serum 25(OH)D concentrations and COVID-19 infectivity and severity. Although the results from randomized clinical trials aiming to prove the benefit of vitamin D supplementation for these purposes are pending, there is no downside to increasing vitamin D intake and having sensible sunlight exposure to maintain serum 25(OH)D at a level of at least 30 ng/mL (75 nmol/L) and preferably at 40 to 60 ng/mL (100–150 nmol/L)<sup>12</sup> to achieve optimal health benefits of vitamin D and minimize the risk of COVID-19 infection and its severity.

## Disclosure

Michael F. Holick is a former consultant for Quest Diagnostics Inc., a consultant for Biogena Inc. and Ontometrics Inc., and on the speaker's Bureau for Abbott Inc.

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## References

- Hu B, Guo H, Zhou P, Shi ZL. Characteristics of SARS-CoV-2 and COVID-19. *Nat Rev Microbiol.* 2020;19(3):141–154.
- The Lancet Infectious Diseases. COVID-19, a pandemic or not? *Lancet Infect Dis.* 2020;20(4):383.
- COVID-19 coronavirus pandemic 2021. <https://www.worldometers.info/coronavirus/#countries>. Accessed February 18, 2021.
- Kordzadeh-Kermani E, Khalili H, Karimzadeh I. Pathogenesis, clinical manifestations and complications of coronavirus disease 2019 (COVID-19). *Future Microbiol.* 2020;15:1287–1305.
- Wiersinga WJ, Rhodes A, Cheng AC, Peacock SJ, Prescott HC. Pathophysiology, transmission, diagnosis, and treatment of coronavirus disease 2019 (COVID-19): a review. *JAMA.* 2020;324(8):782–793.
- Wolff D, Nee S, Hickey NS, Marschollek M. Risk factors for Covid-19 severity and fatality: a structured literature review. *Infection.* 2021;49(1):15–28.
- Snowden LR, Graaf G. COVID-19, social determinants past, present, and future, and African Americans' health. *J Racial Ethn Health Disparities.* 2021;8(1):12–20.
- Khunti K, Singh AK, Pareek M, Hanif W. Is ethnicity linked to incidence or outcomes of covid-19? *BMJ.* 2020;369:m1548.
- Wacker M, Holick MF. Sunlight and vitamin D: a global perspective for health. *Dermatoendocrinol.* 2013;5(1):51–108.
- Charoennangam N, Shirvani A, Holick MF. Vitamin D for skeletal and non-skeletal health: what we should know. *J Clin Orthop Trauma.* 2019;10(6):1082–1093.
- Holick MF. Vitamin D deficiency. *N Engl J Med.* 2007;357(3):266–281.
- Holick MF, Binkley NC, Bischoff-Ferrari HA, et al. Evaluation, treatment, and prevention of vitamin D deficiency: an Endocrine Society clinical practice guideline. *J Clin Endocrinol Metab.* 2011;96(7):1911–1930.
- Charoennangam N, Holick MF. Immunologic effects of vitamin D on human health and disease. *Nutrients.* 2020;12(7):2097.
- Holick MF. High prevalence of vitamin D inadequacy and implications for health. *Mayo Clin Proc.* 2006;81(3):353–373.
- Malabanan A, Veronikis IE, Holick MF. Redefining vitamin D insufficiency. *Lancet.* 1998;351(9105):805–806.
- Thomas MK, Lloyd-Jones DM, Thadhani RI, et al. Hypovitaminosis D in medical inpatients. *N Engl J Med.* 1998;338(12):777–783.
- Holick MF, Siris ES, Binkley N, et al. Prevalence of vitamin D inadequacy among postmenopausal North American women receiving osteoporosis therapy. *J Clin Endocrinol Metab.* 2005;90(6):3215–3224.
- Hewison M. Vitamin D and immune function: an overview. *Proc Nutr Soc.* 2012;71(1):50–61.
- Adams JS, Hewison M. Unexpected actions of vitamin D: new perspectives on the regulation of innate and adaptive immunity. *Nat Clin Pract Endocrinol Metab.* 2008;4(2):80–90.
- Bouillon R, Marcocci C, Carmeliet G, et al. Skeletal and extraskeletal actions of vitamin D: current evidence and outstanding questions. *Endocr Rev.* 2019;40(4):1109–1151.
- Crowe FL, Steur M, Allen NE, Appleby PN, Travis RC, Key TJ. Plasma concentrations of 25-hydroxyvitamin D in meat eaters, fish eaters, vegetarians and vegans: results from the EPIC-Oxford study. *Public Health Nutr.* 2011;14(2):340–346.
- Liu J, Arcot J, Cunningham J, et al. New data for vitamin D in Australian foods of animal origin: impact on estimates of national adult vitamin D intakes in 1995 and 2011–13. *Asia Pac J Clin Nutr.* 2015;24(3):464–471.
- Liu J, Greenfield H, Strobel N, Fraser DR. The influence of latitude on the concentration of vitamin D3 and 25-hydroxy-vitamin D3 in Australian red meat. *Food Chem.* 2013;140(3):432–435.
- Al Mutairi AN, Nasrat GH, Russell DW. Mutation of the CYP2R1 vitamin D 25-hydroxylase in a Saudi Arabian family with severe vitamin D deficiency. *J Clin Endocrinol Metab.* 2012;97(10):E2022–E2025.
- Gombart AF. The vitamin D antimicrobial peptide pathway and its role in protection against infection. *Future Microbiol.* 2009;4(9):1151–1165.
- Aranow C. Vitamin D and the immune system. *J Investig Med.* 2011;59(6):881–886.
- Lemire JM, Archer DC, Beck L, Spiegelberg HL. Immunosuppressive actions of 1,25-dihydroxyvitamin D3: preferential inhibition of Th1 functions. *J Nutr.* 1995;125(suppl 6):S1704–S1708.
- Boonstra A, Barrat FJ, Crain C, Heath VL, Savelkoul HFJ, O'Garra A. 1alpha,25-Dihydroxyvitamin d3 has a direct effect on naive CD4(+) T cells to enhance the development of Th2 cells. *J Immunol.* 2001;167(9):4974–4980.
- Tang J, Zhou R, Luger D, et al. Calcitriol suppresses antiretinal autoimmunity through inhibitory effects on the Th17 effector response. *J Immunol.* 2009;182(8):4624–4632.
- Chen S, Sims GP, Chen XX, Gu YY, Chen S, Lipsky PE. Modulatory effects of 1,25-dihydroxyvitamin D3 on human B cell differentiation. *J Immunol.* 2007;179(3):1634–1647.
- Kongsbak M, Levring TB, Geisler C, von Essen MR. The vitamin D receptor and T cell function. *Front Immunol.* 2013;4:148.
- Sarkar S, Hewison M, Studzinski GP, Li YC, Kalia V. Role of vitamin D in cytotoxic T lymphocyte immunity to pathogens and cancer. *Crit Rev Clin Lab Sci.* 2016;53(2):132–145.
- Shirvani A, Kalajian TA, Song A, Holick MF. Disassociation of vitamin D's calcemic activity and non-calcemic genomic activity and individual responsiveness: a randomized controlled double-blind clinical trial. *Sci Rep.* 2019;9(1):17685.
- Shirvani A, Kalajian TA, Song A, et al. Variable genomic and metabolomic responses to varying doses of vitamin D supplementation. *Anticancer Res.* 2020;40(1):535–543.
- Carlberg C, Seuter S, de Mello VDF, et al. Primary vitamin D target genes allow a categorization of possible benefits of vitamin D<sub>3</sub> supplementation. *PLoS One.* 2013;8(7), e71042.
- Hansdottir S, Monick MM, Hinde SL, Lovan N, Look DC, Hunninghake GW. Respiratory epithelial cells convert inactive vitamin D to its active form: potential effects on host defense. *J Immunol.* 2008;181(10):7090–7099.
- Tripathi S, Tecle T, Verma A, Crouch E, White M, Hartshorn KL. The human cathelicidin LL-37 inhibits influenza A viruses through a mechanism distinct from that of surfactant protein D or defensins. *J Gen Virol.* 2013;94(Pt 1):40–49.
- Sousa FH, Casanova V, Findlay F, et al. Cathelicidins display conserved direct antiviral activity towards rhinovirus. *Peptides.* 2017;95:76–83.
- Barlow PG, Svoboda P, Mackellar A, et al. Antiviral activity and increased host defense against influenza infection elicited by the human cathelicidin LL-37. *PLoS One.* 2011;6(10), e25333.
- Quraishi SA, De Pascale G, Needleman JS, et al. Effect of cholecalciferol supplementation on vitamin D status and cathelicidin levels in sepsis: a randomized, placebo-controlled trial. *Crit Care Med.* 2015;43(9):1928–1937.
- Roth A, Lütke S, Meinberger D, et al. LL-37 fights SARS-CoV-2: the vitamin D-inducible peptide LL-37 inhibits binding of SARS-CoV-2 spike protein to its cellular receptor angiotensin converting enzyme 2 in vitro. *Preprint Posted online December 2, 2020 bioRxiv 408153.* <https://doi.org/10.1101/2020.12.02.408153>.
- Jiang JS, Chou HC, Chen CM. Cathelicidin attenuates hyperoxia-induced lung injury by inhibiting oxidative stress in newborn rats. *Free Radic Biol Med.* 2020;150:23–29.
- Lu L, Zhang H, Dauphars DJ, He YW. A potential role of interleukin 10 in COVID-19 pathogenesis. *Trends Immunol.* 2021;42(1):3–5.
- McElvaney OJ, Hobbs BD, Qiao D, et al. A linear prognostic score based on the ratio of interleukin-6 to interleukin-10 predicts outcomes in COVID-19. *EBioMedicine.* 2020;61:103026.
- Heine G, Niesner U, Chang H-D, et al. 1,25-dihydroxyvitamin D(3) promotes IL-10 production in human B cells. *Eur J Immunol.* 2008;38(8):2210–2218.
- Ashtari F, Toghianifar N, Zarkeesh-Esfahani SH, Mansourian M. Short-term effect of high-dose vitamin D on the level of interleukin 10 in patients with multiple sclerosis: a randomized, double-blind, placebo-controlled clinical trial. *Neuroimmunomodulation.* 2015;22(6):400–404.
- Gopinath K, Danda D. Supplementation of 1,25 dihydroxy vitamin D3 in patients with treatment naïve early rheumatoid arthritis: a randomised controlled trial. *Int J Rheum Dis.* 2011;14(4):332–339.
- Finamor DC, Sinigaglia-Coimbra R, Neves LCM, et al. A pilot study assessing the effect of prolonged administration of high daily doses of vitamin D on the clinical course of vitiligo and psoriasis. *Dermatoendocrinol.* 2013;5(1):222–234.
- Disphanurat W, Viarasilpa W, Chakkavittumrong P, Pongcharoen P. The clinical effect of oral vitamin D2 supplementation on psoriasis: a double-

- blind, randomized, placebo-controlled study. *Dermatol Res Pract.* 2019;2019:5237642.
50. McLaughlin L, Clarke L, Khalilidehkordi E, Butzkeven H, Taylor B, Broadley SA. Vitamin D for the treatment of multiple sclerosis: a meta-analysis. *J Neurol.* 2018;265(12):2893–2905.
  51. Li J, Chen N, Wang D, Zhang J, Gong X. Efficacy of vitamin D in treatment of inflammatory bowel disease: a meta-analysis. *Med (Baltimore).* 2018;97(46):e12662.
  52. Evans RM, Lippman SM. Shining light on the COVID-19 pandemic: A vitamin D receptor checkpoint in defense of unregulated wound healing. *Cell Metab.* 2020;32(5):704–709.
  53. Ajabshir S, Asif A, Nayer A. The effects of vitamin D on the renin-angiotensin system. *J Nephropathol.* 2014;3(2):41–43.
  54. Ali RM, Al-Shorbagy MY, Helmy MW, El-Abhar HS. Role of Wnt4/β-catenin, Ang II/TGFβ, ACE2, NF-κB, and IL-18 in attenuating renal ischemia/reperfusion-induced injury in rats treated with Vit D and pioglitazone. *Eur J Pharmacol.* 2018;831:68–76.
  55. Wu J, Deng W, Li S, Yang X. Advances in research on ACE2 as a receptor for 2019-nCoV. *Cell Mol Life Sci.* 2021;78(2):531–544.
  56. Hanff TC, Harhay MO, Brown TS, Cohen JB, Mohare AM. Is there an association between COVID-19 mortality and the renin-angiotensin system? A call for epidemiologic investigations. *Clin Infect Dis.* 2020;71(15):870–874.
  57. Yuan W, Pan W, Kong J, et al. 1,25-Dihydroxyvitamin D3 suppresses renin gene transcription by blocking the activity of the cyclic AMP response element in the renin gene promoter. *J Biol Chem.* 2007;282(41):29821–29830.
  58. Xu J, Yang J, Chen J, Luo Q, Zhang Q, Zhang H. Vitamin D alleviates lipopolysaccharide-induced acute lung injury via regulation of the renin-angiotensin system. *Mol Med Rep.* 2017;16(5):7432–7438.
  59. Ma TKW, Kam KKH, Yan BP, Lam YY. Renin–angiotensin–aldosterone system blockade for cardiovascular diseases: current status. *Br J Pharmacol.* 2010;160(6):1273–1292.
  60. Garvin MR, Alvarez C, Miller JL, et al. A mechanistic model and therapeutic interventions for COVID-19 involving a RAS-mediated bradykinin storm. *Elife.* 2020;9, e59177.
  61. Gibson CC, Davis CT, Zhu W, et al. Dietary vitamin D and its metabolites non-genomically stabilize the endothelium. *PLoS One.* 2015;10(10), e0140370.
  62. Vila Cuena M, Ferrantelli E, Meinster E, et al. Vitamin D attenuates endothelial dysfunction in uremic rats and maintains human endothelial stability. *J Am Heart Assoc.* 2018;7(17), e008776.
  63. Mohammad S, Mishra A, Ashraf MZ. Emerging role of vitamin D and its associated molecules in pathways related to pathogenesis of thrombosis. *Biomolecules.* 2019;9(11):649.
  64. Rolf L, Muris A-H, Hupperts R, Damoiseaux J. Illuminating vitamin D effects on B cells—the multiple sclerosis perspective. *Immunology.* 2016;147(3):275–284.
  65. Yamamoto EA, Nguyen JK, Liu J, et al. Low levels of vitamin D promote memory B cells in lupus. *Nutrients.* 2020;12(2):291.
  66. Hope-Simpson RE. The role of season in the epidemiology of influenza. *J Hyg (Lond).* 1981;86(1):35–47.
  67. Li Y, Wang X, Nair H. Global seasonality of human seasonal coronaviruses: a clue for postpandemic circulating season of severe acute respiratory syndrome coronavirus 2? *J Infect Dis.* 2020;222(7):1090–1097.
  68. Ianevski A, Zusinaite E, Shtaida N, et al. Low temperature and low UV indexes correlated with peaks of influenza virus activity in Northern Europe during 2010–2018. *Viruses.* 2019;11(3):207.
  69. Shaman J, Jeon CY, Giovannucci E, Lipsitch M. Shortcomings of vitamin D-based model simulations of seasonal influenza. *PLoS One.* 2011;6(6), e20743.
  70. Tamerius JD, Shaman J, Alonso WJ, et al. Environmental predictors of seasonal influenza epidemics across temperate and tropical climates. *PLoS Pathog.* 2013;9(3), e1003194.
  71. Cannell JJ, Vieth R, Umhau JC, et al. Epidemic influenza and vitamin D. *Epidemiol Infect.* 2006;134(6):1129–1140.
  72. Sabetta JR, DePetrillo P, Cipriani RJ, Smardin J, Burns LA, Landry ML. Serum 25-hydroxyvitamin D and the incidence of acute viral respiratory tract infections in healthy adults. *PLoS One.* 2010;5(6), e11088.
  73. Ingham TR, Jones B, Camargo CA, et al. Association of vitamin D deficiency with severity of acute respiratory infection: A case-control study in New Zealand children. *Eur Respir J.* 2014;44(suppl 58):439.
  74. Urashima M, Segawa T, Okazaki M, Kurihara M, Wada Y, Ida H. Randomized trial of vitamin D supplementation to prevent seasonal influenza A in schoolchildren. *Am J Clin Nutr.* 2010;91(5):1255–1260.
  75. Martineau AR, Jolliffe DA, Hooper RL, et al. Vitamin D supplementation to prevent acute respiratory tract infections: systematic review and meta-analysis of individual participant data. *BMJ.* 2017;356:i6583.
  76. Rello J, Valenzuela-Sánchez F, Ruiz-Rodriguez M, Moyano S. Sepsis: a review of advances in management. *Adv Ther.* 2017;34(11):2393–2411.
  77. de Haan K, Groeneveld ABJ, de Geus HRH, Egal M, Struijs A. Vitamin D deficiency as a risk factor for infection, sepsis and mortality in the critically ill: systematic review and meta-analysis. *Crit Care.* 2014;18(6):660.
  78. Vipul P, Shuchi C, Avinash A, Manish G, Sukriti K, Ved P. Correlation of serum vitamin D level with mortality in patients with sepsis. *Indian J Crit Care Med.* 2017;21(4):199–204.
  79. Rübsamen D, Kunze MM, Buderus V, et al. Inflammatory conditions induce IRES-dependent translation of cyp24a1. *PLoS One.* 2014;9(1), e85314.
  80. Dahl B, Schiødt FV, Ott P, et al. Plasma concentration of Gc-globulin is associated with organ dysfunction and sepsis after injury. *Crit Care Med.* 2003;31(1):152–156.
  81. Han JE, Jones JL, Tangpricha V, et al. High dose vitamin D administration in ventilated intensive care unit patients: a pilot double blind randomized controlled trial. *J Clin Transl Endocrinol.* 2016;4:59–65.
  82. Martucci G, McNally D, Parekh D, et al. Trying to identify who may benefit most from future vitamin D intervention trials: a post hoc analysis from the VITDAL-ICU study excluding the early deaths. *Crit Care.* 2019;23(1):200.
  83. National Heart, Lung, and Blood Institute PETAL Clinical Trials Network, Ginde AA, Brower RG, et al. Early high-dose vitamin D3 for critically ill, vitamin D-deficient patients. *N Engl J Med.* 2019;381(26):2529–2540.
  84. Annweiler G, Corvaisier M, Gautier J, et al. Vitamin D supplementation associated to better survival in hospitalized frail elderly COVID-19 patients: the GERIA-COVID quasi-experimental study. *Nutrients.* 2020;12(11).
  85. Annweiler C, Hanotte B, Grandin de l'Eprevier C, Sabatier JM, Lafaa L, Celarier T. Vitamin D and survival in COVID-19 patients: a quasi-experimental study. *J Steroid Biochem Mol Biol.* 2020;204:105771.
  86. Kaufman HW, Niles JK, Kroll MH, Bi C, Holick MF. SARS-CoV-2 positivity rates associated with circulating 25-hydroxyvitamin D levels. *PLoS One.* 2020;15(9), e0239252.
  87. Melitzer DO, Best TJ, Zhang H, Vokes T, Arora V, Solway J. Association of vitamin D status and other clinical characteristics with COVID-19 test results. *JAMA Netw Open.* 2020;3(9), e2019722.
  88. Im JH, Je YS, Baek J, Chung M-H, Kwon HY, Lee J-S. Nutritional status of patients with COVID-19. *Int J Infect Dis.* 2020;100:390–393.
  89. Merzon E, Tworowski D, Gorohovski A, et al. Low plasma 25(OH) vitamin D level is associated with increased risk of COVID-19 infection: an Israeli population-based study. *FEBS Journal.* 2020;287(17):3693–3702.
  90. Hernández JL, Nan D, Fernandez-Ayala M, et al. Vitamin D status in hospitalized patients with SARS-CoV-2 infection. *J Clin Endocrinol Metab.* 2021;106(3):e1343–e1353.
  91. Ferrari D, Locatelli M. No significant association between vitamin D and COVID-19. A retrospective study from a northern Italian hospital. *Int J Vitam Nutr Res.* 2020;1–4.
  92. Hastie CE, Mackay DF, Ho F, et al. Vitamin D concentrations and COVID-19 infection in UK Biobank. *Diabetes Metab Syndr.* 2020;14(4):561–565.
  93. Pereira M, Dantas Damascena A, Galvão Azevedo LM, de Almeida Oliveira T, da Mota Santana J. Vitamin D deficiency aggravates COVID-19: systematic review and meta-analysis. *Crit Rev Food Sci Nutr.* 2020;1–9.
  94. Hars M, Mendes A, Serrattice C, et al. Sex-specific association between vitamin D deficiency and COVID-19 mortality in older patients. *Osteoporos Int.* 2020;31(12):2495–2496.
  95. Walk J, Dofferhoff ASM, van den Ouwerland JMW, van Daal H, Janssen R. Vitamin D – contrary to vitamin K – does not associate with clinical outcome in hospitalized COVID-19 patients. Preprint. Posted online November 9, 2020. medRxiv 20227512. <https://doi.org/10.1101/2020.11.07.20227512>
  96. Karahan S, Katkat F. Impact of serum 25(OH) vitamin D level on mortality in patients with COVID-19 in Turkey. *J Nutr Health Aging.* 2021;25(2):189–196.
  97. Szeto B, Zucker JE, LaSota ED, et al. Vitamin D status and COVID-19 clinical outcomes in hospitalized patients. *Endocr Res.* 2020;1–8.
  98. Mendi A, Apewokin S, Wells AA, Morrow AL. Factors associated with hospitalization and disease severity in a racially and ethnically diverse population of COVID-19 patients. *medRxiv : the preprint server for health sciences.* 2020, 2020.06.25.20137323.
  99. Radujkovic A, Hippchen T, Tiwari-Heckler S, Dreher S, Boxberger M, Merle U. Vitamin D deficiency and outcome of COVID-19 patients. *Nutrients.* 2020;12(9):2757.
  100. Ling SF, Broad E, Murphy R, et al. High-dose cholecalciferol booster therapy is associated with a reduced risk of mortality in patients with COVID-19: a cross-sectional multi-centre observational study. *Nutrients.* 2020;12(12):3799.
  101. Rejnmark L, Bislev LS, Cashman KD, et al. Non-skeletal health effects of vitamin D supplementation: a systematic review on findings from meta-analyses summarizing trial data. *PLoS One.* 2017;12(7), e0180512.
  102. Vranic L, Mikolašević I, Milić S. Vitamin D deficiency: consequence or cause of obesity? *Medicina (Kaunas).* 2019;55(9):541.
  103. Levin A, Bakris GL, Molitch M, et al. Prevalence of abnormal serum vitamin D, PTH, calcium, and phosphorus in patients with chronic kidney disease: results of the study to evaluate early kidney disease. *Kidney Int.* 2007;71(1):31–38.
  104. Maghbooli Z, Sahraian MA, Ebrahimi M, et al. Vitamin D sufficiency, a serum 25-hydroxyvitamin D at least 30–ng/ml, reduced risk for adverse clinical outcomes in patients with COVID-19 infection. *PLoS One.* 2020;15(9), e0239799.
  105. French CB, McDonnell SL, Vieth R. 25-Hydroxyvitamin D variability within-person due to diurnal rhythm and illness: a case report. *J Med Case Rep.* 2019;13(1):29.
  106. Smolders J, van den Ouwerland J, Geven C, Pickkers P, Kox M. Letter to the editor: vitamin D deficiency in COVID-19: mixing up cause and consequence. *Metabolism.* 2021;115:154434.

107. Baktash V, Hosack T, Patel N, et al. Vitamin D status and outcomes for hospitalised older patients with COVID-19. *Postgrad Med J.* 2020; postgradmedj-2020-138712.
108. Entrenas Castillo M, Entrenas Costa LM, Vaquero Barrios JM, et al. Effect of calcifediol treatment and best available therapy versus best available therapy on intensive care unit admission and mortality among patients hospitalized for COVID-19: a pilot randomized clinical study. *J Steroid Biochem Mol Biol.* 2020;203:105751.
109. Murai IH, Fernandes AL, Sales LP, et al. Effect of a single high dose of vitamin D3 on hospital length of stay in patients with moderate to severe COVID-19: a randomized clinical trial. *JAMA.* 2021.
110. Holick MF. Resurrection of vitamin D deficiency and rickets. *J Clin Invest.* 2006;116(8):2062–2072.
111. Di Filippo L, Formenti AM, Rovere-Querini P, et al. Hypocalcemia is highly prevalent and predicts hospitalization in patients with COVID-19. *Endocrine.* 2020;68(3):475–478.
112. Luxwolda MF, Kuipers RS, Kema IP, van der Veer E, Dijck-Brouwer DAJ, Muskiet FAJ. Vitamin D status indicators in indigenous populations in East Africa. *Eur J Nutr.* 2013;52(3):1115–1125.
113. Holick MF. The death D-fying vitamin. *Mayo Clin Proc.* 2018;93(6):679–681.
114. Charoenngam N, Shirvani A, Holick MF. The ongoing D-lemma of vitamin D supplementation for nonskeletal health and bone health. *Curr Opin Endocrinol Diabetes Obes.* 2019;26(6):301–305.
115. Dudenkov DV, Mara KC, Pettersson TM, Maxson JA, Thacher TD. Serum 25-hydroxyvitamin D values and risk of all-cause and cause-specific mortality: a population-based cohort study. *Mayo Clin Proc.* 2018;93(6):721–730.
116. Hosseini-nezhad A, Holick MF. Vitamin D for health: a global perspective. *Mayo Clin Proc.* 2013;88(7):720–755.
117. Ekwari JP, Zwicker JD, Holick MF, Giovannucci E, Veugelers PJ. The importance of body weight for the dose response relationship of oral vitamin D supplementation and serum 25-hydroxyvitamin D in healthy volunteers. *PLoS One.* 2014;9(11):e111265.
118. Cooper ID, Crofts CAP, DiNicolantonio JJ, et al. Relationships between hyperinsulinaemia, magnesium, vitamin D, thrombosis and COVID-19: rationale for clinical management. *Open Heart.* 2020;7(2), e001356.
119. Dai Q, Zhu X, Manson JE, et al. Magnesium status and supplementation influence vitamin D status and metabolism: results from a randomized trial. *Am J Clin Nutr.* 2018;108(6):1249–1258.